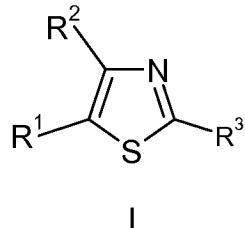


In the Claims:

The current status of all claims is listed below and supersedes all previous lists of claims.

Please cancel claims 7-11 without prejudice to their presentation in another application, and amend claims 1-3 as follows:

1. (currently amended) A compound of formula (I)



wherein

R^1 and R^2 are independently selected from phenyl, thiaryl, and pyridyl, each of which is independently optionally substituted with one, two or three Z groups;

Z is selected from a C_{1-6} alkyl group, a C_{1-6} alkoxy group, hydroxy, halo, trifluoromethyl, trifluoromethylthio, trifluoromethoxy, trifluoromethylsulphonyl, nitro, amino, mono or di C_{1-3} alkylamino, mono or di C_{1-3} alkylamido, C_{1-3} alkylsulphonyl, C_{1-3} alkoxycarbonyl, carboxy, cyano, carbamoyl, mono or di C_{1-3} alkyl carbamoyl, sulphamoyl, acetyl, $-\text{O}-\text{CH}_2-\text{CH}_2-\text{O}-$ attached at two adjacent carbons, and phenyl, optionally substituted with one or more of the following: a C_{1-6} alkyl group, trifluoromethyl, a C_{1-6} alkoxy group, trifluoromethoxy, halo, or $-\text{O}-\text{CH}_2-\text{CH}_2-\text{O}-$ attached at two adjacent carbons;

R^3 is $-\text{X}-\text{Y}-\text{NR}^4\text{R}^5$;

R^4 and R^5 are independently selected from:

~~a C_{1-6} alkyl group, optionally substituted with a C_{1-6} alkoxy group or trifluoromethoxy;~~

~~an (amino) C_{1-4} alkyl group, wherein the amino is optionally substituted by one or more C_{1-3} alkyl groups;~~

~~a non-aromatic C_{3-15} carboyclic group, optionally substituted with a C_{1-3} alkoxy C_{1-3} alkyl group;~~

a ~~(C₃₋₁₂cycloalkyl)C₁₋₃alkyl~~ group;

a ~~(CH₂)_r(phenyl)_s~~ group, wherein r is 0, 1, 2, 3 or 4, and wherein s is 1 when r is 0, otherwise s is 1 or 2, and wherein the phenyl groups are optionally independently substituted with one, two or three Z groups;

naphthyl;

anthraenyl;

a saturated 5-to-8 membered heterocyclic piperidine group containing one nitrogen and optionally containing one of the following: oxygen, sulphur or an additional nitrogen, wherein the heterocyclic piperidine group is optionally substituted by one or more C₁₋₃alkyl groups or benzyl;

1-adamantylmethyl; and

a -(CH₂)_tHet group, wherein t is 0, 1, 2, 3 or 4, and the alkylene chain is optionally substituted by one or more C₁₋₃alkyl groups, and wherein Het is ~~an aromatic heterocycle optionally substituted by one, two or three groups selected from a C₁₋₆alkyl group; a C₁₋₆alkoxy group, trifluoromethoxy or halo~~ or Het is a saturated 5-to-8 membered heterocyclic piperidine group containing one nitrogen and optionally containing one of the following: oxygen, sulphur or an additional nitrogen; wherein the heterocyclic piperidine group is optionally substituted by one or more C₁₋₃alkyl groups, hydroxy or benzyl; and

wherein R⁴ may be H; and

wherein R⁴ and R⁵ taken together with the nitrogen atom to which they are attached form a saturated 5-to-8 membered heterocyclic piperidine group containing one nitrogen and optionally containing one of the following: oxygen, sulphur or an additional nitrogen; wherein the heterocyclic piperidine group is optionally substituted with one or more C₁₋₃alkyl groups, hydroxy or benzyl;

X is CO or SO₂; and

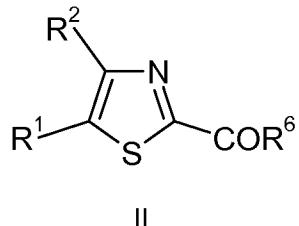
Y is absent or represents NH optionally substituted by a C₁₋₃alkyl group;

or a pharmaceutically acceptable salt, prodrug or solvate thereof;

with the proviso that R¹ and R² are not both 4-methoxyphenyl and the proviso that when

R^1 is phenyl and R^2 represents phenyl or 4-fluorophenyl, X is CO and Y is absent then the group NR^4R^5 is not methyl-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]amino, methylpiperazino, 2-[1-methyl-4-piperidinyl]ethylamino, or [2-[1-(phenylmethyl)-4-piperidinyl]ethyl]amino.

2. (currently amended) A compound of formula I as represented by formula (II)



wherein

R^1 is phenyl, optionally substituted by one or more of the following: a C₁₋₆alkyl group, trifluoromethyl, a C₁₋₆alkoxy group, trifluoromethoxy, halo, or -O-CH₂-CH₂-O- attached at two adjacent carbons;

R^2 is phenyl, optionally substituted by one or more of the following: a C₁₋₆alkyl group, trifluoromethyl, a C₁₋₆alkoxy group, trifluoromethoxy, halo, or -O-CH₂-CH₂-O- attached at two adjacent carbons; and

R^6 is selected from 1-piperidinylamino, a ~~C₃₋₇cycloalkylamino group, optionally substituted by C₁₋₆alkoxyC₁₋₂alkyl, pyridylamino, wherein the pyridyl ring is optionally substituted by one or more of the following: a C₁₋₆alkyl group; a C₁₋₆alkoxy group or trifluoromethoxy; a C₁₋₆alkylamino group, wherein the alkyl chain is optionally substituted by one or more of the following: a C₁₋₆alkoxy group, trifluoromethoxy or morpholino;~~ or a pharmaceutically acceptable salt, prodrug or solvate thereof;

~~with the proviso that when R¹ is 4-methoxyphenyl and R² is 4-methoxyphenyl, then R⁶ is not 2-(morpholino)ethyl.~~

3. (currently amended) A compound selected from:

~~4-(4-chlorophenyl)-5-(2,4-dichlorophenyl)thiazole-2-carboxylic acid cyclohexylamide;~~
~~5-(4-chlorophenyl)-4-(2,4-dichlorophenyl)thiazole-2-carboxylic acid cyclohexylamide;~~
~~4-(4-chlorophenyl)-5-(2,4-dichlorophenyl)thiazole-2-carboxylic acid piperidin-1-~~

ylamide;

5-(4-chlorophenyl)-4-(2,4-dichlorophenyl)thiazole-2-carboxylic acid piperidin-1-ylamide;

~~4-(4-bromophenyl)-5-phenylthiazole-2-carboxylic acid cyclohexylamide;~~

~~4-(4-bromophenyl)-5-phenylthiazole-2-carboxylic acid piperidin-1-ylamide;~~

~~4,5-bis(4-chlorophenyl)thiazole-2-carboxylic acid cyclohexylamide;~~

~~4,5-bis(4-chlorophenyl)thiazole-2-carboxylic acid piperidin-1-ylamide;~~

~~4-(4-methoxyphenyl)-5-phenylthiazole-2-carboxylic acid cyclohexylamide;~~

~~4,5-bis(4-methoxyphenyl)thiazole-2-carboxylic acid cyclohexylamide;~~

~~4,5-bis(4-methoxyphenyl)thiazole-2-carboxylic acid piperidin-1-ylamide;~~

~~5-(7-bromo-2,3-dihydrobenzo[1,4]dioxin-6-yl)-4-phenylthiazole-2-carboxylic acid piperidin-1-ylamide;~~

~~4-(7-bromo-2,3-dihydrobenzo[1,4]dioxin-6-yl)-5-phenylthiazole-2-carboxylic acid piperidin-1-ylamide;~~

~~4,5-bis(4-chlorophenyl)thiazole-2-carboxylic acid (2-methoxymethylcyclopentyl)-amide;~~

~~4,5-bis(4-chlorophenyl)thiazole-2-carboxylic acid pyridin-4-ylamide;~~

~~4,5-bis(4-chlorophenyl)thiazole-2-carboxylic acid (2-ethoxyethyl)amide; and~~

~~4,5-bis(4-chlorophenyl)thiazole-2-carboxylic acid (2-morpholin-4-yl-ethyl)amide~~

and where applicable, optical isomers, tautomers, stereoisomers and racemates thereof as well as pharmaceutically acceptable salts ~~and solvates~~ thereof.

4. (canceled).

5. (previously presented) A pharmaceutical formulation comprising a compound of any one of claims 1 to 3 and a pharmaceutically acceptable adjuvant, diluent or carrier.

6-11. (canceled).

REMARKS